## **Amendments to the Claims:**

This listing of claims will replace all prior versions, and listings, of claims in the application:

## What is claimed is:

1. (Currently amended) A process for preparing a chloropurine compound of formula (I)

er a derivative thereof, which comprises ring closure of a compound of formula (VII) or a derivative thereof

(VII)

in the presence of <u>a</u> catalytic <u>amount of</u> acid and at least one equivalent of a <del>formate</del> derivative trialkylorthoformate.

- 2. (Original) A process according to claim 1 wherein the acid is sulfuric acid, hydrochloric acid, or an alkyl or arylsulfonic acid.
- 3. (Previously presented) A process according to claim 1 wherein the acid is present in an amount of from 0.05 to 0.1 equivalents by mole based on an amount of the compound of formula (VII).
- 4. (Currently amended) A process according to claim 1 wherein the formate derivative trialkylorthoformate is triethylorthoformate.

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5. (Currently amended) A process according claim 1 wherein the formate derivative trialkylorthoformate is present in an amount of 1 to 1.5 equivalents by mole based on the amount of the compound of formula (VII).

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6. (Currently amended) A process according to claim 1 wherein the compound of formula (VII) er a derivative thereof is prepared by condensing an amino alcohol of formula (IV) er a derivative thereof

(IV)

with a compound of formula (VIII) or a derivative thereof

(VIII)

in the presence of a base.

- 7. (Original) A process according to claim 6 wherein the condensation reaction is carried out in n-butanol in the presence of sodium bicarbonate.
- 8. (Original) A process according to claim 6 wherein the condensation reaction is carried out in n-butanol in the presence of anhydrous potassium carbonate.
- 9. (Currently amended) A process according to claim 1 wherein the chloropurine compound of formula (I) or derivative thereof prepared by the ring closure reaction is converted *in situ* to abacavir or a derivative thereof.
- 10. Cancelled